

L8 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:675777 CAPLUS <<LOGINID::20080113>>
 DOCUMENT NUMBER: 147:64481
 TITLE: Methods of screening agents for activity using teleosts
 INVENTOR(S): McGrath, Patricia; Parng, Chuenlei; Serbedzija, George N.
 PATENT ASSIGNEE(S): Phylonix Pharmaceuticals, Inc., USA
 SOURCE: U.S. Pat. Appl. Publ., 74pp., Cont.-in-part of U.S. Ser. No. 678,765.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2007143865	A1	20070621	US 2006-515355	20060901
US 6299858	B1	20011009	US 1999-255397	19990222 <--
EP 1548123	A1	20050629	EP 2005-1508	19990222 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY				
US 6656449	B1	20031202	US 2000-645432	20000823 <--
US 2004062712	A1	20040401	US 2003-678765	20031002 <--
US 2006104905	A1	20060518	US 2005-280849	20051115
PRIORITY APPLN. INFO.:				
			US 1998-75783P	P 19980223
			US 1998-100950P	P 19980918
			US 1999-255397	A2 19990222
			US 2000-645432	A1 20000823
			US 2003-678765	A2 20031002
			EP 1999-934309	A3 19990222
			US 2001-947635	A1 20010905

ABSTRACT:
 The present invention provides methods of screening an agent for activity using teleosts. Methods of screening an agent for angiogenesis activity, toxic activity and an effect cell death activity in teleosts are provided. Methods of screening an agent including a drug for an activity in the brain or central nervous system in zebrafish are provided. The invention further provides high throughput methods of screening agents in multi-well plates. Functional blood-brain barrier (BBB) was present in embryo zebrafish demonstrating it's usefulness for screening agents for BBB permeability.

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2007143865	A1	20070621	US 2006-515355	20060901
US 6299858	B1	20011009	US 1999-255397	19990222 <--
EP 1548123	A1	20050629	EP 2005-1508	19990222 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY				
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US 2004062712	A1	20040401	US 2003-678765	20031002 <--
US 2006104905	A1	20060518	US 2005-280849	20051115
IT 50-02-2, Dexamethasone	302-79-4, Retinoic acid	23110-15-8D,		
Fumagillin, derivative	629627-77-6, Ovicillin			
RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); BIOL (Biological study)				
(methods of screening agents for activity using teleosts)				
IT 50-81-7, Ascorbic acid, biological studies	53-06-5, Cortisone	59-02-9,		
α -Tocopherol	70-51-9, Deferoxamine	91-56-5, Isatin	348-67-4,	

D-Methionine 452-86-8, 4-Methylcatechol 616-91-1, N-Acetyl-L-cysteine
700-06-1, Indole-3-carbinol 2323-36-6, Deprenyl 2682-49-7,
2-Oxothiazolidine 10118-90-8, Minocycline 27138-57-4 50903-99-6,
L-NAME 57828-26-9, Lipoic acid
RL: PAC (Pharmacological activity); BIOL (Biological study)
(methods of screening agents for activity using teleosts)

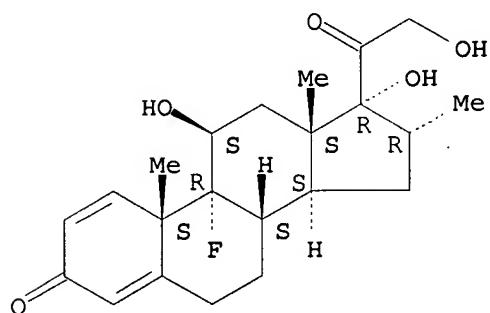
IT 50-02-2, Dexamethasone

RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); BIOL (Biological study)
(methods of screening agents for activity using teleosts)

RN 50-02-2 CAPLUS

CN Pregna-1,4-diene-3,20-dione, 9-fluoro-11,17,21-trihydroxy-16-methyl-,
(11 β ,16 α)- (CA INDEX NAME)

Absolute stereochemistry.

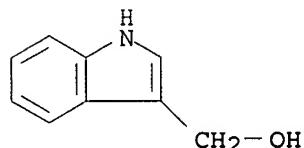


IT 700-06-1, Indole-3-carbinol

RL: PAC (Pharmacological activity); BIOL (Biological study)
(methods of screening agents for activity using teleosts)

RN 700-06-1 CAPLUS

CN 1H-Indole-3-methanol (CA INDEX NAME)



L8 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:376660 CAPLUS <<LOGINID::20080113>>

DOCUMENT NUMBER: 138:379207

TITLE: Improved use of antitumoral compound in cancer therapy

INVENTOR(S): Jimeno, Jose; Ruiz Casado, Ana; Lopez Lazaro, Luis;
Rowinsky, Eric; Hidalgo, Manuel

PATENT ASSIGNEE(S): Pharmamar S.A., Spain

SOURCE: PCT Int. Appl., 28 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.

KIND DATE

APPLICATION NO.

DATE

WO 2003039571	A1	20030515	WO 2002-US33548	20021021 <--
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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CN 1606449	A	20050413	CN 2002-825666	20021021 <--
JP 2005509650	T	20050414	JP 2003-541862	20021021 <--
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MX 2004PA03674	A	20040723	MX 2004-PA3674	20040419
IN 2004DN01070	A	20060728	IN 2004-DN1070	20040421
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US 2005004018	A1	20050106	US 2004-492320	20040818
PRIORITY APPLN. INFO.:			US 2001-348414P	P 20011019
			WO 2002-US33548	W 20021021

ABSTRACT:

Improved dosing schedules for ecteinascidin 743 are given for treatment of cancer.

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003039571	A1	20030515	WO 2002-US33548	20021021 <--
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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IN 2004DN01070	A	20060728	IN 2004-DN1070	20040421
NO 2004002035	A	20040518	NO 2004-2035	20040518
US 2005004018	A1	20050106	US 2004-492320	20040818
IT 114899-77-3, Ecteinascidin 743				

RL: PKT (Pharmacokinetics); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(improved use of antitumoral compound in cancer therapy)

IT 50-02-2 15663-27-1, Cisplatin 23214-92-8, Doxorubicin
33069-62-4, Paclitaxel

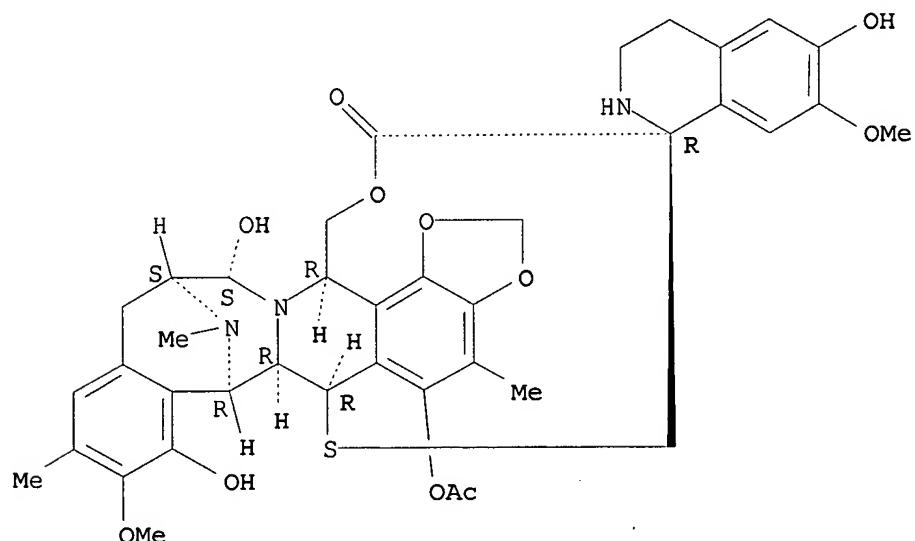
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(improved use of antitumoral compound in cancer therapy)

IT 114899-77-3, Ecteinascidin 743
RL: PKT (Pharmacokinetics); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(improved use of antitumoral compound in cancer therapy)

RN 114899-77-3 CAPLUS

CN Spiro[6,16-(epithiopropanoxymethano)-7,13-imino-12H-1,3-dioxolo[7,8]isoquinol[3,2-b][3]benzazocine-20,1'(2'H)-isoquinolin]-19-one,
5-(acetoxy)-3',4',6,6a,7,13,14,16-octahydro-6',8,14-trihydroxy-7',9-dimethoxy-4,10,23-trimethyl-, (1'R,6R,6aR,7R,13S,14S,16R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



IT 50-02-2

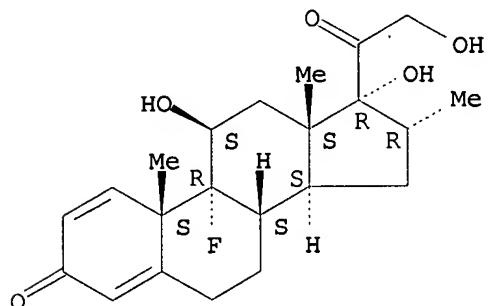
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(improved use of antitumoral compound in cancer therapy)

RN 50-02-2 CAPLUS

CN Pregna-1,4-diene-3,20-dione, 9-fluoro-11,17,21-trihydroxy-16-methyl-, (11 β ,16 α)- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2002:353292 CAPLUS <<LOGINID::20080113>>
DOCUMENT NUMBER: 136:350548
TITLE: Effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions
INVENTOR(S): Takahashi, Naoto; Weitman, Steve; D'incalci, Maurizio; Faicloth, Glynn Thomas; Giavazzi, Rafaella; Gescher, Andreas
PATENT ASSIGNEE(S): Pharma Mar, S.A., Spain
SOURCE: PCT Int. Appl., 26 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002036135	A2	20020510	WO 2001-GB4902	20011106 <--
WO 2002036135	A3	20030410		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2428160	A1	20020510	CA 2001-2428160	20011106 <--
AU 200212499	A	20020515	AU 2002-12499	20011106 <--
BR 2001015162	A	20031021	BR 2001-15162	20011106 <--
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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
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NZ 525730	A	20041224	NZ 2001-525730	20011106 <--
NO 2003002027	A	20030704	NO 2003-2027	20030506 <--
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ZA 2003003474	A	20040806	ZA 2003-3474	20030506 <--
BG 107843	A	20040630	BG 2003-107843	20030523 <--
US 2004108086	A1	20040610	US 2003-416086	20030917 <--
PRIORITY APPLN. INFO.:			US 2000-246233P	P 20001106
			US 2000-248095P	P 20001113
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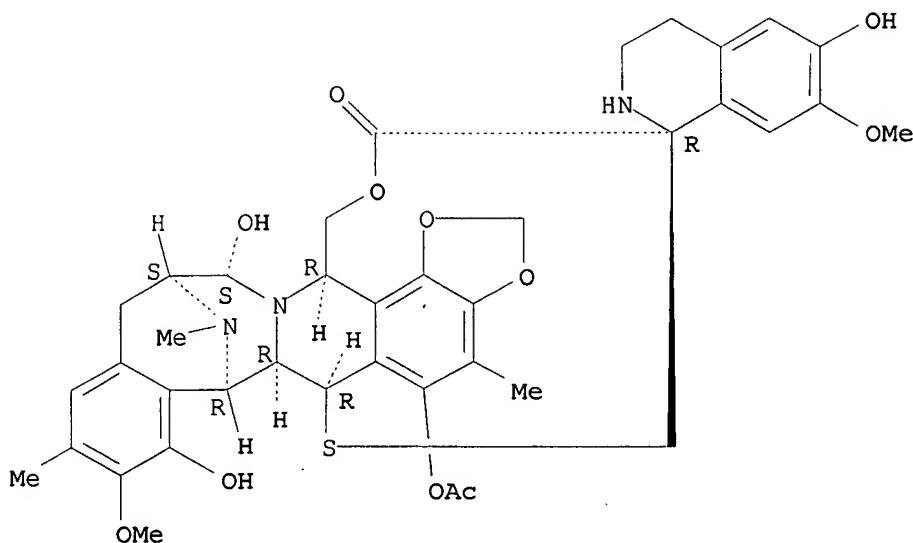
ABSTRACT:

ET-743 is used in the preparation of a medicament for an effective treatment of a tumor by combination therapy employing ET-743 with another drug.

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2002036135	A2	20020510	WO 2001-GB4902	20011106 <--

WO 2002036135	A3	20030410		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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EP 1365808	A2	20031203	EP 2001-980710	20011106 <--
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HU 2004000648	A2	20040628	HU 2004-648	20011106 <--
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BG 107843	A	20040630	BG 2003-107843	20030523 <--
US 2004108086	A1	20040610	US 2003-416086	20030917 <--
IT <u>114899-77-3, ET-743</u>	RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions)			
IT <u>50-02-2, Dexamethasone</u>	RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hepatoprotectant; effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions)			
IT <u>114899-77-3, ET-743</u>	RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions)			
RN 114899-77-3 CAPLUS				
CN Spiro[6,16-(epithiopropanoxymethano)-7,13-imino-12H-1,3-dioxolo[7,8]isoquino[3,2-b][3]benzazocine-20,1'(2'H)-isoquinolin]-19-one, 5-(acetyloxy)-3',4',6,6a,7,13,14,16-octahydro-6',8,14-trihydroxy-7',9-dimethoxy-4,10,23-trimethyl-, (1'R,6R,6aR,7R,13S,14S,16R)- (CA INDEX NAME)				

Absolute stereochemistry. Rotation (-).



IT 50-02-2, Dexamethasone

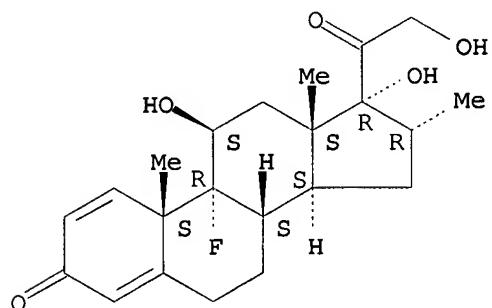
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(hepatoprotectant; effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions)

RN 50-02-2 CAPLUS

CN Pregna-1,4-diene-3,20-dione, 9-fluoro-11,17,21-trihydroxy-16-methyl-, (11 β ,16 α)- (CA INDEX NAME)

Absolute stereochemistry.



L8 ANSWER 4 OF 5 USPATFULL on STN

ACCESSION NUMBER: 2005:4912 USPATFULL <<LOGINID::20080113>>

TITLE: Use of antitumoral compound in cancer therapy

INVENTOR(S): Jimeno, Jose, Madrid, SPAIN

Casado, Ana Ruiz, Madrid, SPAIN

Lazaro, Luis Lopez, Madrid, SPAIN

Rowensky, Eric, San Antonio, TX, UNITED STATES

Hidalgo, Manuel, Baltimore, MD, UNITED STATES

NUMBER	KIND	DATE
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PATENT INFORMATION: US 2005004018 A1 20050106

APPLICATION INFO.: US 2004-492320 A1 20040818 (10) <--

NUMBER DATE

PRIORITY INFORMATION: US 2001-348414P 20011019 (60)
 DOCUMENT TYPE: Utility
 FILE SEGMENT: APPLICATION
 LEGAL REPRESENTATIVE: FISH & RICHARDSON PC, 225 FRANKLIN ST, BOSTON, MA,
 02110
 NUMBER OF CLAIMS: 19
 EXEMPLARY CLAIM: 1
 LINE COUNT: 675
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 ABSTRACT:

Improved dosing schedules for ecteinascidin 743 are given for treatment of cancer.

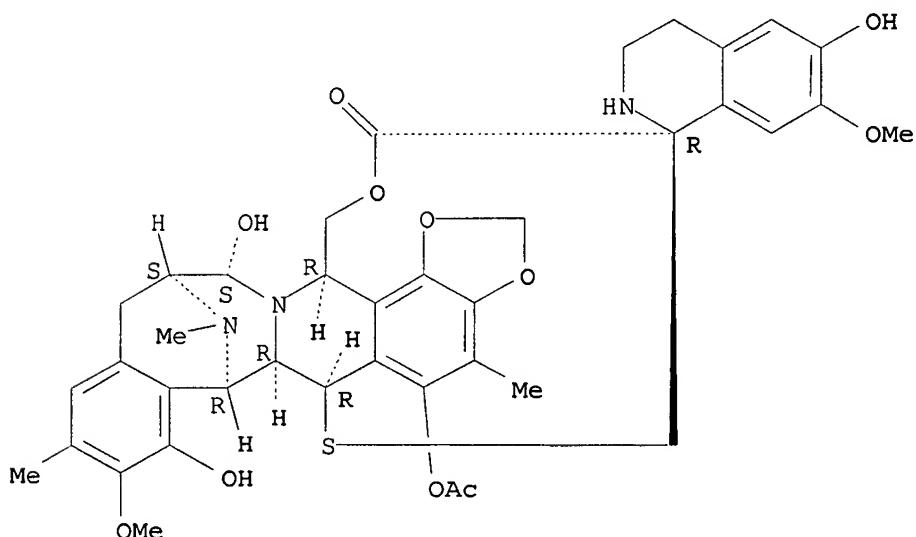
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AI US 2004-492320 A1 20040818 (10)
 20021021

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IT 114899-77-3, Ecteinascidin 743
 (improved use of antitumoral compound in cancer therapy)
 IT 50-02-2 15663-27-1, Cisplatin 23214-92-8, Doxorubicin
 33069-62-4, Paclitaxel
 (improved use of antitumoral compound in cancer therapy)
 IT 114899-77-3, Ecteinascidin 743
 (improved use of antitumoral compound in cancer therapy)
 RN 114899-77-3 USPATFULL
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 one, 5-(acetyloxy)-3',4',6,6a,7,13,14,16-octahydro-6',8,14-trihydroxy-
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Absolute stereochemistry. Rotation (-).



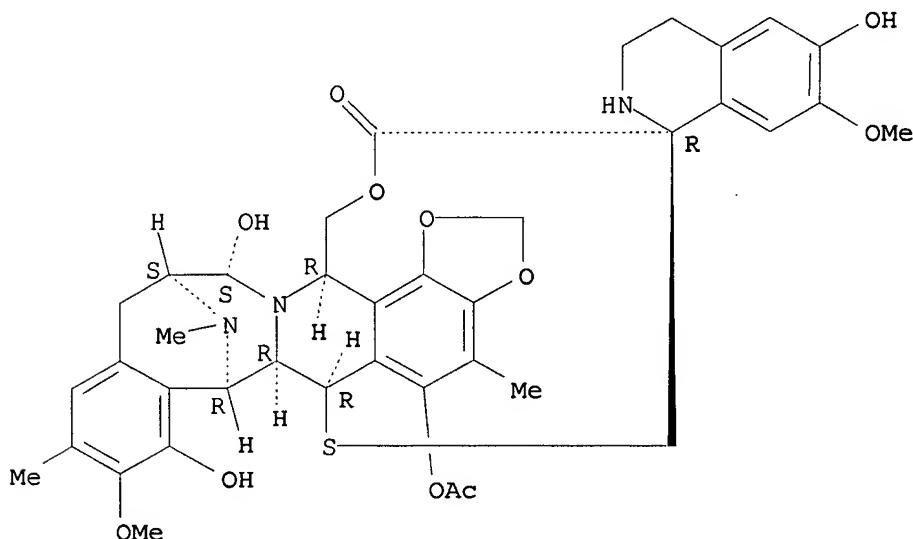
IT 50-02-2
 (improved use of antitumoral compound in cancer therapy)

antitumor agents resulting in additive and synergistic interactions)

RN 114899-77-3 USPATFULL

CN Spiro[6,16-(epithiopropanoxymethano)-7,13-imino-12H-1,3-dioxolo[7,8]isoquinol[3,2-b][3]benzazocine-20,1' (2'H)-isoquinolin]-19-one, 5-(acetyloxy)-3',4',6,6a,7,13,14,16-octahydro-6',8,14-trihydroxy-7',9-dimethoxy-4,10,23-trimethyl-, (1'R,6R,6aR,7R,13S,14S,16R)- (CA INDEX NAME)

Absolute stereochemistry. Rotation (-).



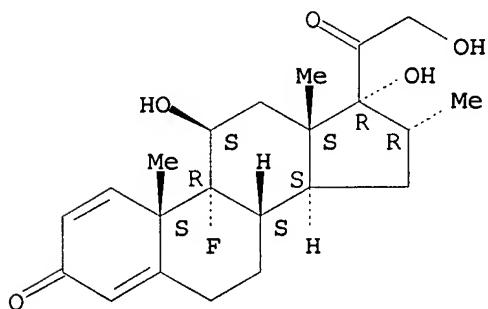
IT 50-02-2, Dexamethasone

(hepatoprotectant; effective antitumor treatments with combinations of ET-743 with other antitumor agents resulting in additive and synergistic interactions)

RN 50-02-2 USPATFULL

CN Pregna-1,4-diene-3,20-dione, 9-fluoro-11,17,21-trihydroxy-16-methyl-,
(11 β ,16 α)- (CA INDEX NAME)

Absolute stereochemistry.



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E DEXAMETHASONE/CN
L2 1 S E15
E INDOLE-3-CARBINOL/CN
L3 1 S E27

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ON 13 JAN 2008

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